16 June 2000

Article reference: CB14.160600 Coffee Break archives

nature Story contributed by Alison Mitchell, Nature Reviews Molecular Cell Biology



The Drosophila protein EAST is involved in the assembly of a nucleoskeleton that may prevent chromosome collisions.

Click here for more information.

## The beginning of the END

What do traditional office skills like filing and book-keeping have in common with cell biology? The answer lies in the organization of essential components - be they documents or proteins - into distinct compartments. Take the nucleus, for example, where the proteins involved in processes such as transcription and RNA metabolism are physically clustered, and chromosomes are partitioned into discrete territories.

Reporting in Nature Cell Biology, Martin Wasser and William Chia from the National University of Singapore describe a molecular secretary that may organize this nuclear filing system in the fruit fly Drosophila melanogaster. Known as EAST (for 'enhanced adult sensory threshold'), it contains 12 potential proteolytic sites as well as seven putative nuclear-localization signals. These characteristics led the authors to propose that EAST is an unstable protein targeted specifically to the nucleus.

To test this prediction, Wasser and Chia studied the expression pattern of EAST in giant nuclei from Drosophila salivary glands. Confocal microscopy captured images of EAST in a region that the authors refer to as the extrachromosomal nuclear domain (END), which, as its name suggests, is the area around and between the chromosomes. Similar compartments have previously been detected in other organisms, where they are thought to belong to a putative nuclear endoskeleton.

What might be the function of EAST within this 'nucleoskeleton'? The authors reasoned that, if it is involved in forming a structural backbone, EAST may recruit other factors to the END. So they studied the effect of disrupting the east gene on distribution of a protein called CP60 that normally co-localizes with EAST. Consistent with their ideas, the CP60 expression pattern was destroyed. What's more, Wasser and Chia found overlapping expression patterns between EAST and nuclear actin - a satisfying discovery given that, outside the nucleus, actin is a core structural component of the cytoskeleton.

Overexpression of EAST gave equally striking results - an expansion of the END. This effect, which can also be induced by heat-shocking Drosophila, results from accumulation of the extra EAST. But as the figure shows, the consequences of expansion vary depending where the END expands: if EAST accumulates mainly between the chromosomes (left), the effect will be to increase the spacing between them; expansion between the chromosomes and nuclear membrane (right), however, will compress the chromosomes. In either case, this could prevent the random collision of neighbouring chromosome arms when cells are under stress.

Wasser and Chia have shown, then, that EAST is a nuclear architect involved in the assembly of an expandable nucleoskeleton between chromosomes. The next steps will be to work out how increased levels of EAST cause the END to expand, and to take a closer look at how the EAST protein is regulated.

#### Live PubMed searches

|| The nucleoskeleton || REVIEWS on the nuclear matrix |

#### Genome tutorial

|| Find CP60 in the Drosophila genome ||

#### Further NCBI resources

|| CP60 in LocusLink || Drosophila homepage ||

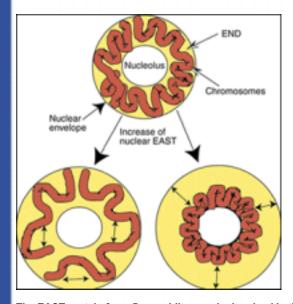
Comments? Questions? We would welcome feedback on NCBI's Coffee Break. Email to: info@ncbi.nlm.nih.gov



# S NCBI Figure Coffee Break

Coffee Break Article Genome tutorial

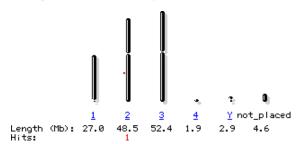
Genome Links Genomic Biology homepage Genomes FTP site Map viewer help



The EAST protein from Drosophila may be involved in the assembly of a nucleoskeleton. When overexpressed, EAST expands the extrachromosomal nuclear domain (END) - the space between chromosomes. The consequences of expansion vary depending where the END expands: if EAST accumulates mainly between the chromosomes (left), the effect will be to increase the spacing between them; expansion between the chromosomes and nuclear membrane (right), however, will compress the chromosomes. In either case, this could prevent the random collision of neighbouring chromosome arms when cells are under stress.



#### **Drosophila melanogaster** genome view



#### **Precomputed BLAST results:**

Enter any protein ID (accession or gi number) to find matches to *Drosophila* genome proteins



Search results for query "CP60": 1 hit

All data displayed on these pages are based on the sequence provided by Celera and the Berkeley Drosophila Genome Project. Searching this sequence for CP60 shows one hit, localized on chromosome 2, in the dynamically drawn figure (above).

The table below lists the hits, and the corresponding sequence information, if any. In this case, the Celera/Berkeley genome has been annotated to include CP60, so the sequence view (sv), genbank view (gv), etc, of this segment is available.

Click on the Map60 link in the Map Element column to view the MAP60 gene within its genomic context.



#### Links:

Map element - Map view

Nucleotide:

sv Sequence Viewer

**gb** GenBank view of record

gr GenBank view of region

fr FASTA format of region

Protein:

gp GenPept view

fp FASTA protein format

Resources:

b Precomputed BLAST

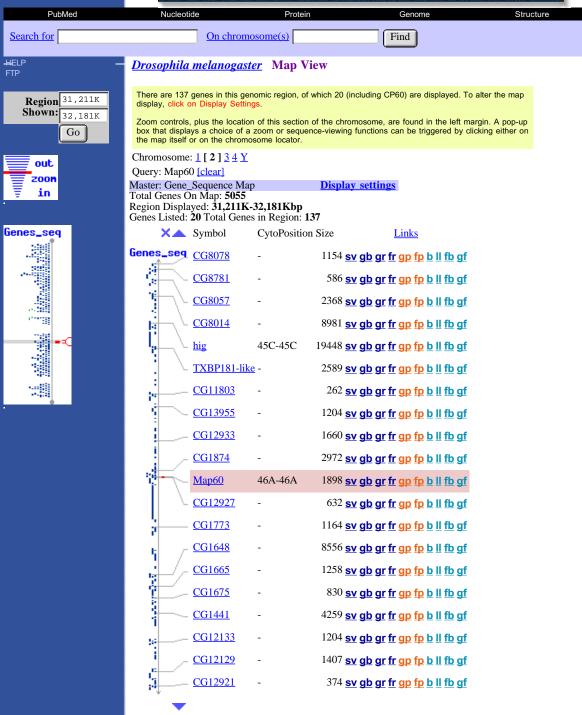
| LocusLink

**fb** FlyBase

**gf** GadFly

Disclaimer | Write to the Help Desk NCBI | NLM | NIH





Disclaimer | Write to the Help Desk NCBI | NLM | NIH



Find

PubMed Nucleotide Protein Genome Structure Taxonomy

Genomic Biology

Search for

CTD alta

About this sequence

Sequencing Centers

Calars

Related Resources

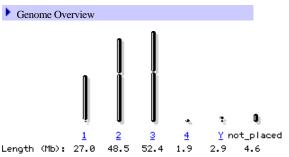
FlvBase

GadFlv

The Interactive Fly

#### Drosophila melanogaster genome

On chromosome(s)





The assembled and annotated genome sequence of the euchromatic arms of the five *Drosophila melanogaster* (fruit fly) chromosomes is now available in GenBank. The sequence, determined in a collaboration between <u>Celera</u> and the <u>Berkeley Drosophila Genome Project</u>, is described in the March 24, 2000 issue of <u>Science</u>. The ~137 Mb of sequence, most of which is found on

chromosomes 1 (also known as X), 2, and 3, contains ~13,500 annotated genes. ~2470 of these genes correlate with a known gene described in FlyBase. FlyBase provides sequence for an additional ~500 genes that are not annotated on the Celera/BDGP sequence.

From early observations of the banding patterns of its polytene chromosomes to current work on mRNA and protein gradients in the developing embryo, *Drosophila melanogaster* has been studied in biology labs for over eighty years. Many of the genes that define the spatial pattern of cell types and body parts have now been identified, along with the regulatory pathways in which they operate. As a number of these genes have counterparts in higher eukaryotes, the study of the *Drosophila* developmental program provides insight into human development as well. *Drosophila* is the second multicellular organism to be sequenced, after the nematode *Caenorhabditis elegans*.

### Resources

- ▶ Get sequence data by FTP
- Entrez Map Viewer [About...]
  Genome view
  Map View
  Sequence view
- Search for genes in <u>LocusLink</u>
- ▶ Related Structures:

Drosophila genome proteins with sequence similarity to proteins with known structure

- BLAST your sequence against the *Drosophila melanogaster* genome
- Precomputed BLAST results:
  [More...]

Enter the gene or protein ID of a Drosophila genome sequence protein to view protein matches in all organisms



Enter the ID (accession or gi number) of a protein from any organism to view matches to *Drosophila* genome sequence proteins

Any Protein ID	
	Show

Questions or Comments?
Write to the NCBI Service Desk

Revised May 9, 2000